

chain nodes:

7 8 9 10 11 12 18 27 28 29 31

ring nodes:

1 2 3 4 5 6 21 22 23 24 25 26

chain bonds:

6-8 7-23 8-18 10-11 11-12 18-28 26-27 27-29

ring bonds:

1-2 1-6 2-3 3-4 4-5 5-6 21-22 21-26 22-23 23-24 24-25 25-26

exact/norm bonds:

6-8 7-23 8-18 18-28 26-27 27-29

exact bonds:

10-11 11-12 21-22 21-26 22-23 23-24 24-25 25-26

normalized bonds:

1-2 1-6 2-3 3-4 4-5 5-6

isolated ring systems:

containing 1: 21:

G1:[\*1-\*2],[\*3-\*4]

Match level:

1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:CLASS8:CLASS9:CLASS10:CLASS11:CLASS12:CLASS18:CLASS 20:Atom 21:Atom 22:Atom 23:Atom 24:Atom 25:Atom 26:Atom 27:CLASS28:Atom 29:Atom 31:Atom 32:Atom

Generic attributes:

28:

Saturation

: Unsaturated

29:

Saturation 31: Saturation

: Unsaturated

: Unsaturated

=> Uploading C:\Program Files\Stnexp\Queries\10517594.str

```
chain nodes :
7 8 9 10 11 12 18 27 28 29 31
ring nodes :
1 2 3 4 5 6 21 22 23 24 25 26
chain bonds :
6-8 7-23 8-18 10-11 11-12 18-28 26-27 27-29
ring bonds :
1-2 1-6 2-3 3-4 4-5 5-6 21-22 21-26 22-23 23-24 24-25 25-26
exact/norm bonds :
6-8 7-23 8-18 18-28 26-27 27-29
exact bonds :
10-11 11-12 21-22 21-26 22-23 23-24 24-25 25-26
normalized bonds :
1-2 1-6 2-3 3-4 4-5 5-6
isolated ring systems :
containing 1 : 21 :
```

G1:[\*1-\*2],[\*3-\*4]

Match level:
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:CLASS 8:CLASS 9:CLASS 10:CLASS 11:CLASS 12:CLASS 18:CLASS 20:Atom 21:Atom 22:Atom 23:Atom 24:Atom 25:Atom 26:Atom 27:CLASS 28:Atom 29:Atom 31:Atom 32:Atom Generic attributes:

28:

Saturation : Unsaturated

29:

Saturation : Unsaturated

31:

Saturation : Unsaturated

L1 STRUCTURE UPLOADED

=> d 11

L1 HAS NO ANSWERS

L1 STR

 $Ak^{1}$ 

Structure attributes must be viewed using STN Express query preparation.

=> s 11 sss sam
SAMPLE SEARCH INITIATED 16:33:09 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 6392 TO ITERATE

31.3% PROCESSED 2000 ITERATIONS
INCOMPLETE SEARCH (SYSTEM LIMIT EXCEEDED)
SEARCH TIME: 00.00.01

O ANSWERS

FULL FILE PROJECTIONS: O

ONLINE \*\*COMPLETE\*\*

BATCH \*\*COMPLETE\*\*

10/517,594

PROJECTED ITERATIONS: 123047 TO 132633 PROJECTED ANSWERS: 0 TO 0

0 SEA SSS SAM L1

24 SEA SSS FUL L1

=> s ll sss ful

FULL SEARCH INITIATED 16:33:34 FILE 'REGISTRY'

FULL SCREEN SEARCH COMPLETED - 130540 TO ITERATE

100.0% PROCESSED 130540 ITERATIONS

SEARCH TIME: 00.00.04

24 ANSWERS

=> => s 13 L4

1 L3

=> d 14 bib, ab, hitstr

```
ANSWER 1 OF 1 CAPLUS COPYRIGHT 2007 ACS on STN
L4
     2003:1006977 CAPLUS
AN
     140:59654
DN
     Preparation of substituted diaminopyrimidines as protein kinase C \theta
TI
     inhibitors
     Baudler, Monika; Bhagwat, Shripad S.; Erdman, Paul E.; Baudler, Monika;
IN
     Bhagwat, Shripad S.; Erdman, Paul E.; Gantner, Florian; Palanki, Moorthy
     S. S.; Schudt, Christian; Stadlwieser, Josef; Zapf, James
     Altana Pharma A.-G., Germany; Signal Pharmaceuticals, Inc.
PA
SO
     PCT Int. Appl., 42 pp.
     CODEN: PIXXD2
DT
     Patent
     English
LA:
FAN.CNT 1
                                             APPLICATION NO.
                                                                     DATE
     PATENT NO.
                         KIND
                                 DATE
                                             _____
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                                 _____
     _____
                                                                    20030607
                                             WO 2003-EP6016
                          `A1
                                 20031224
PΙ
             AE, AL, AU, BA, BR, CA, CN, CO, CU, DZ, EC, GE, HR, ID, IL, IN,
             IS, JP, KR, LT, LV, MA, MK, MX, NO, NZ, PH, PL, SG, TN, UA, US,
             VN, YU, ZA, ZW
         RW: AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE,
             DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE,
             SI, SK, TR
                                             CA 2003-2489458
                                                                     20030607
                                 20031224
     CA 2489458
                           Α1
                                                                     20030607
                                 20031231
                                             AU 2003-236720
     AU 2003236720
                           Α1
                                                                     20030607
                                 20050323
                                             EP 2003-735581
     EP 1515964
                           Α1
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             IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK
                                                                     20041213
                          Α1
                                 20051006
                                             US 2004-517594
   · US 2005222186
PRAI US 2002-388312P
                           Ρ
                                 20020614
                                 20020620
     EP 2002-13675
                          Α
                                 20030607
                          W
     WO 2003-EP6016
OS
     MARPAT 140:59654
     Pyrimidinediamines I [R1-R3 = (un)substituted mono- or bicyclic aromatic; R4,
AB
     R5 = H, Me; A1, A2 = alkylene, CH2CH2O] were prepared for use as protein
     kinase C \theta inhibitors. Thus, 4-amino-1-benzylpiperidine was treated
     with 5-bromo-2,4-dichloropyrimidine, followed by 2-aminomethylpyridine to
     give {5-bromo-2-[(2-pyridylmethyl)amino]pyrimidin-4-yl}(1-benzyl-4-
     piperidinyl)amine which was treated with 3-thiopheneboronic acid to give I
     [A1, A2 = CH2, R1 = 3-thienyl, R2 = 2-pyridyl, R3 = Ph, R4, R5 = H] which
     had an IC50 for inhibition of protein kinase C \theta < 4.0 \muM.
     637356-54-8P 637356-68-4P 637356-72-0P
ΙT
     637356-78-6P 637356-79-7P 637356-80-0P
     637356-81-1P 637356-82-2P 637356-83-3P
     637356-84-4P 637356-85-5P 637356-86-6P
     637356-87-7P 637356-89-9P 637356-90-2P
     637356-91-3P
     RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU
     (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES
         (preparation of substituted diaminopyrimidines as protein kinase C \boldsymbol{\theta}
        inhibitors)
     637356-54-8 CAPLUS
RN
     2,4-Pyrimidinediamine, N4-[1-(phenylmethyl)-4-piperidinyl]-N2-(2-
CN
     pyridinylmethyl)-5-(3-thienyl)- (9CI) (CA INDEX NAME)
```

RN 637356-68-4 CAPLUS
CN Benzamide, 4-[4-[[1-(phenylmethyl)-4-piperidinyl]amino]-2-[(4-pyridinylmethyl)amino]-5-pyrimidinyl]- (9CI) (CA INDEX NAME)

RN 637356-72-0 CAPLUS
CN 2,4-Pyrimidinediamine, N4-[1-(phenylmethyl)-4-piperidinyl]-N2-(4-pyridinylmethyl)-5-(2-thienyl)- (9CI) (CA INDEX NAME)

RN 637356-78-6 CAPLUS
CN 2,4-Pyrimidinediamine, 5-(2-furanyl)-N4-[1-(phenylmethyl)-4-piperidinyl]N2-(2-pyridinylmethyl)- (9CI) (CA INDEX NAME)

RN 637356-79-7 CAPLUS
CN 2,4-Pyrimidinediamine, 5-(3-furanyl)-N4-[1-(phenylmethyl)-4-piperidinyl]N2-(2-pyridinylmethyl)- (9CI) (CA INDEX NAME)

637356-80-0 CAPLUS RN CN

2,4-Pyrimidinediamine, N4-[1-(phenylmethyl)-4-piperidinyl]-N2-(2pyridinylmethyl)-5-(2-thienyl)- (9CI) (CA INDEX NAME)

637356-81-1 CAPLUS RN

Benzamide, 4-[4-[[1-(phenylmethyl)-4-piperidinyl]amino]-2-[(2-pyridinylmethyl)amino]-5-pyrimidinyl]- (9CI) (CA INDEX NAME) CN

RN 637356-82-2 CAPLUS

CN 2,4-Pyrimidinediamine, 5-(3-chloro-4-fluorophenyl)-N4-[1-(phenylmethyl)-4-piperidinyl]-N2-(2-pyridinylmethyl)- (9CI) (CA INDEX NAME)

RN 637356-83-3 CAPLUS

CN Acetamide, N-[3-[4-[[1-(phenylmethyl)-4-piperidinyl]amino]-2-[(2-pyridinylmethyl)amino]-5-pyrimidinyl]phenyl]- (9CI) (CA INDEX NAME)

RN 637356-84-4 CAPLUS

CN 2,4-Pyrimidinediamine, 5-(1,3-benzodioxol-5-yl)-N4-[1-(phenylmethyl)-4-piperidinyl]-N2-(2-pyridinylmethyl)- (9CI) (CA INDEX NAME)

RN 637356-85-5 CAPLUS

CN Phenol, 3-[4-[[1-(phenylmethyl)-4-piperidinyl]amino]-2-[(2-pyridinylmethyl)amino]-5-pyrimidinyl]- (9CI) (CA INDEX NAME)

RN 637356-86-6 CAPLUS

CN Phenol, 4-[4-[[1-(phenylmethyl)-4-piperidinyl]amino]-2-[(2-pyridinylmethyl)amino]-5-pyrimidinyl]- (9CI) (CA INDEX NAME)

RN 637356-87-7 CAPLUS

CN Ethanone, 1-[4-[4-[[1-(phenylmethyl)-4-piperidinyl]amino]-2-[(2-pyridinylmethyl)amino]-5-pyrimidinyl]phenyl]- (9CI) (CA INDEX NAME)

RN 637356-89-9 CAPLUS

CN Ethanone, 1-[3-[4-[[1-(phenylmethyl)-4-piperidinyl]amino]-2-[(2-pyridinylmethyl)amino]-5-pyrimidinyl]phenyl]- (9CI) (CA INDEX NAME)

RN 637356-90-2 CAPLUS

CN Benzamide, N,N-dimethyl-4-[4-[[1-(phenylmethyl)-4-piperidinyl]amino]-2-[(2-pyridinylmethyl)amino]-5-pyrimidinyl]- (9CI) (CA INDEX NAME)

RN 637356-91-3 CAPLUS
CN Acetamide, N-[4-[4-[[1-(phenylmethyl)-4-piperidinyl]amino]-2-[(2-pyridinylmethyl)amino]-5-pyrimidinyl]phenyl]- (9CI) (CA INDEX NAME)

RN 637356-60-6 CAPLUS
CN 2,4-Pyrimidinediamine, 5-phenyl-N4-[1-(phenylmethyl)-4-piperidinyl]-N2-(4-pyridinylmethyl)- (9CI) (CA INDEX NAME)

RN 637356-63-9 CAPLUS
CN 2,4-Pyrimidinediamine, 5-(4-chlorophenyl)-N4-[1-(phenylmethyl)-4-piperidinyl]-N2-(4-pyridinylmethyl)- (9CI) (CA INDEX NAME)

RN 637356-66-2 CAPLUS

CN 2,4-Pyrimidinediamine, 5-[4-(dimethylamino)phenyl]-N4-[1-(phenylmethyl)-4-piperidinyl]-N2-(4-pyridinylmethyl)- (9CI) (CA INDEX NAME)

RN 637356-70-8 CAPLUS

CN Benzoic acid, 4-[4-[[1-(phenylmethyl)-4-piperidinyl]amino]-2-[(4-pyridinylmethyl)amino]-5-pyrimidinyl]- (9CI) (CA INDEX NAME)

RN 637356-74-2 CAPLUS

CN 2,4-Pyrimidinediamine, 5-(2-furanyl)-N4-[1-(phenylmethyl)-4-piperidinyl]-N2-(4-pyridinylmethyl)- (9CI) (CA INDEX NAME)

RN 637356-75-3 CAPLUS

CN 2,4-Pyrimidinediamine, 5-(3-furanyl)-N4-[1-(phenylmethyl)-4-piperidinyl]-N2-(4-pyridinylmethyl)- (9CI) (CA INDEX NAME)

RN 637356-76-4 CAPLUS
CN 2,4-Pyrimidinediamine, N4-[1-(phenylmethyl)-4-piperidinyl]-N2-(4-pyridinylmethyl)-5-(3-thienyl)- (9CI) (CA INDEX NAME)

RE.CNT 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

## 10/517,594

=> log y COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	5.74	178.50
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	-0.78	-0.78

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